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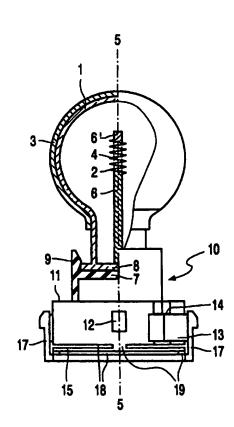
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(54) Title: ELECTRIC LAMP



(57) Abstract: The electric lamp comprises a lamp vessel (1) which is transparent to visible light and which accommodates a light source. At least a part of the lamp vessel (1) is covered with a light-absorbing coating (3). According to the invention, said light-absorbing coating (3) comprises a network which can be obtained by conversion of an organically modified silane by means of a sol-gel process. The organically modified silane is selected from the group formed by compounds of structural formula $R^{1}Si(OR^{11})_{3}$, wherein R^{1} is an alkyl or aryl group and R^{11} is an alkyl group. Preferably, R^{1} is CH_{3} or $C_{0}H_{5}$ and R^{11} is CH_{3} or $C_{2}H_{5}$. Nano-sized silica particles having a diameter $d \le 50$ nm may be incorporated in the network. The pigment is preferably chosen from the group formed by $Fe_{2}O_{3}$, P-doped $Fe_{2}O_{3}$, $ZnFe_{2}O_{4}$, $ZnO.Fe_{2}O_{4}$, $CoAl_{2}O_{4}$, $Nd_{2}O_{5}$, $BiVO_{4}$ and zirconium praseodymium silicate or mixtures thereof. The light-absorbing coating (3) of the electric lamp according to the invention is optically transparent, substantially free of scattering and stable at temperatures up to $350\,^{\circ}C$.



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Electric lamp.

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The invention relates to an electric lamp comprising a light-transmitting lamp vessel which accommodates a light source,

wherein at least a part of the lamp vessel is provided with a light-absorbing coating,

wherein the light-absorbing coating includes a pigment which absorbs a part of the visible light.

Such electric lamps are predominantly used as indicator lamps in vehicles, for example as an amber-colored light source in indicators or as a red-colored light source in brake lights of automobiles. Alternative embodiments of such lamps, wherein the color temperature is increased by means of a light-absorbing coating, can also be used as headlamps of a vehicle. Said light-absorbing coatings are also used as a color layer on (incandescent) lamps for general lighting purposes. Said electric lamps can also be used in traffic lights.

An electric lamp of the type mentioned in the opening paragraph is known from CA-A 0 766 196. In the case of the known electric lamp, a coating is applied to the lamp vessel, which coating comprises a substance which absorbs visible light, for example a dye and/or a pigment.

For the application of said coatings, use is generally made of organic lacquers. The organic lacquer forms a kind of carrier matrix containing the pigment or the dye. Said organic lacquer enables, inter alia, a good adhesion of the coating to the lamp vessel to be obtained. In the known lamp, use is made of a polymethylmethacrylate polymer, which is applied to the lamp vessel by means of dip coating. In an alternative embodiment, a lacquer of a polyester silicone is applied to the lamp vessel by means of a spraying process.

Moreover, use is often made of organic dyes, such as a dye called Zapon 157. Such dyes are added to the lacquer layer to obtain the desired color point.

It is a drawback of the known electric lamp comprising a light-absorbing coating on the basis of an organic lacquer that the adhesion of the coating to the lamp vessel deteriorates substantially and/or the organic dye degrades at temperatures above approximately 220°C. At temperatures close to or higher than said temperature, there is an

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increased risk that the coating cracks and/or becomes detached from the lamp vessel. Since the dimensions of the luminaires accommodating the electric lamp decrease continuously as do the dimensions of the electric lamp itself, the temperature of the lamp vessel provided with the coating currently reaches a temperature of 250°C. In addition, there is a trend towards further miniaturization, so that the lamp vessel provided with the light-absorbing coating reaches temperatures of approximately 325 °C.

It is an object of the invention to provide an electric lamp of the type mentioned in the opening paragraph, wherein the above drawbacks have been obviated.

In accordance with the invention, the electric lamp of the type described in the opening paragraph is characterized in that

the light-absorbing coating comprises a network which can be obtained by conversion of an organically modified silane by means of a sol-gel process,

said organically modified silane being selected from the group formed by compounds of the following structural formula: R^ISi(OR^{II})₃,

wherein R^I comprises an alkyl group or an aryl group, and wherein R^{II} comprises an alkyl group.

By replacing the organic lacquer in the light-absorbing layer in the known electric lamp by a network comprising an organically modified silane as the starting material, an optically transparent, non-scattering, light-absorbing coating is obtained which can resist temperatures up to 400 °C. By using an organically modified silane in the manufacture of the network, a part of the R^I groups, i.e. the alkyl or aryl groups, remain present as an end group in the network. As a result, instead of four network bonds per Si atom, the network in accordance with the invention has fewer than four network bonds per Si atom. This results, by way of example, in a network comprising, on average, approximately three network bonds per Si atom. Despite the fact that the network is partly composed of said alkyl or aryl groups, a network is obtained whose density is at least substantially equal to that of the customary silica network. Unlike the customarily used silica network, a network which is partly composed of said alkyl or aryl groups has a greater elasticity and flexibility. This enables relatively thick light-absorbing coatings to be manufactured.

Preferably, the R^I group comprises CH₃ or C₆H₅. These substances have a relatively good thermal stability. A network comprising methyl or phenyl groups enables thicker coatings to be obtained. Experiments have further shown that coatings wherein

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methyl or phenyl groups are incorporated in a network are stable up to a temperature of at least 350°C. Said groups are end groups in the network and remain part of the network at said higher temperatures. At such a relatively high temperature load on the light-absorbing coating, no appreciable degradation of the network occurs during the service life of the electric lamp.

Preferably, the R^{II} group comprises CH₃ or C₂H₅. Methyl and ethyl groups are particularly suitable because methanol and ethanol are formed in the hydrolysis process, which substances are compatible with the pigment dispersion and evaporate relatively easily. In general, the methoxy groups (-OCH₃) react more rapidly than the ethoxy groups (-OC₂H₅) which, in turn, react more rapidly than (iso)propoxy groups (-OC₃H₇). For a smooth hydrolysis process, use is advantageously made of R^{II} groups which are not very long.

Very suitable starting materials for the manufacture of the network in accordance with the invention are methyltrimethoxy silane (MTMS), where $R^I = R^{II} = CH_3$, methyltriethoxy silane (MTES), where $R^I = CH_3$ and $R^{II} = C_2H_5$, phenyltrimethoxy silane (PTMS), where $R^I = C_6H_5$ and $R^{II} = CH_3$, and phenyltriethoxy silane (PTES), where $R^I = C_6H_5$ and $R^{II} = C_2H_5$. Such starting materials are known per se and commercially available.

An embodiment of the electric lamp in accordance with the invention is characterized in that the thickness t_c of the light-absorbing coating is $t_c \ge 1~\mu m$. If use is made of a network composed of silica, which comprises four network bonds per Si atom, the thickness of the coating is limited, under atmospheric conditions, to approximately at most 0.5 μm . In such silica layers whose thickness exceeds said thickness, stress in the layer readily leads to cracks and/or the coating readily becomes detached from the lamp vessel. By using a network comprising fewer than four network bonds per Si atom, a much thicker layer thickness can be attained. Preferably, $t_c \ge 2~\mu m$. In thicker, light-absorbing coatings, more pigment or dye can be incorporated, whereby the color effect of the coating is improved.

Inorganic filling materials may be incorporated in the light-absorbing coating. For this purpose, in a favorable embodiment of the electric lamp in accordance with the invention, silica particles having a diameter $d \le 50$ nm are incorporated in the network. Incorporation of these so-called nano-sized silica particles reduces shrinkage of the layer during the manufacture thereof. In addition, the incorporation of said nano-sized silica particles makes it possible to obtain even thicker coatings which bond well to the lamp vessel. By adding nano-sized silica particles to a network, wherein alkyl or aryl groups,

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which form the R^I groups, are present as the end group, 20 µm thick layers having favorable bonding properties can be obtained. Such thick layers can contain considerable quantities of a pigment or a dye to obtain the desired color point of the light-absorbing coating. By incorporating said silica particles it becomes possible to manufacture light-absorbing coatings in a larger thickness. The refractive index of such a coating is less influenced by the refractive index of the pigment when the same quantity of pigment is incorporated in a thicker coating. The use of said silica particles thus results in a certain degree of freedom to bring the refractive index of the light-absorbing coating to a desired value and maintain the refractive index at said value.

To manufacture light-absorbing coatings having the desired optical properties, said coatings having the desired thermal stability during the service life of the electric lamp, use is preferably made of inorganic pigments. In a favorable embodiment of the electric lamp in accordance with the invention, the pigment is selected from the group formed by iron oxide, iron oxide doped with phosphor, zinc-iron oxide, cobalt aluminate, neodymium oxide, bismuth vanadate, zirconium praseodymium silicate or mixtures thereof. Iron oxide (Fe₂O₃) is an orange pigment and P-doped Fe₂O₃ is an orange-red pigment. Zinc-iron oxide, for example ZnFe₂O₄ or ZnO.ZnFe₂O₄ are yellow pigments. Mixing (P-doped) Fe₂O₃ with ZnFe₂O₄ yields a pigment of a deep orange color. Cobalt aluminate (CoAl₂O₄) and neodymium oxide (Nd₂O₅) are blue pigments. Bismuth vanadate (BiVO₄), also referred to as pucherite, is a yellow-green pigment. Zirconium praseodymium silicate is a yellow pigment. Experiments have shown that a network including said inorganic pigments does not appreciably degrade during the service life and at the relatively high temperature load on the light-absorbing coating.

In an alternative embodiment, light-absorbing coatings are obtained wherein organic pigments are used. Particularly suitable pigments are the so-called Red 177 (anthraquinone) or chromium phthalic yellow (2RLP) from "Ciba". Further suitable pigments are Red 149 (perylene), Red 122 (quinacridone), Red 257 (Ni-isoindoline), Violet 19 (quinacridone), Blue 15:1 (Cu-phthalocyanine), Green 7 (hal.Cu-phthalocyanine) or Yellow 83 (dyaryl) from "Clariant". Also mixtures of inorganic and organic pigments are suitable, for example a mixture of chromium phthalic yellow and (zinc)iron oxide.

Preferably, an average diameter d_p of the pigment particles is $d_p \le 100$ nm. By using pigments of such relatively small dimensions, optically transparent coatings are obtained which exhibit relatively little light scattering. Since the electric lamp in accordance with the invention is often applied in specially designed reflectors, wherein the light source is

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embodied so as to be punctiform, light scattering by the light-absorbing coatings is an undesirable property. The effect of light scattering is at least substantially precluded if the average diameter of the pigment particles $d_p \le 50$ nm.

In the literature, networks obtained by conversion of an organically modified silane are customarily used to manufacture light-scattering coatings. In this invention, however, the network is used, in particular, to manufacture transparent coatings exhibiting relatively little light scattering.

Particularly suitable electric lamps are obtained by applying a pigment in a light-absorbing coating, which pigment is composed of a mixture of iron oxide and bismuth vanadate, or of a mixture of iron oxide doped with phosphor and bismuth vanadate. Since bismuth vanadate often is only available in a particle size d_p , where $d_p > 100$ nm, a light-absorbing coating comprising such a pigment often exhibits a disturbing degree of light scattering. Inventors have found in experiments that the use of a combination of (P-doped) iron oxide and bismuth vanadate as the pigment causes the light scattering of the coating obtained to be reduced considerably as if the diameter of the bismuth vanadate particles is much smaller than 100 nm. Without being obliged to give any theoretical explanation, the decrease of the light scattering of such a coating is attributed to an increase of the refractive index of the network as a result of the presence of the iron oxide particles.

It has been found that an electric lamp comprising a lamp vessel which is coated in accordance with the invention with a light-absorbing coating comprising a network obtained by conversion of an organically modified silane by means of a sol-gel process preserves its initial properties to a substantial degree during the service life of the electric lamp.

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These and other aspects of the invention will be apparent from and elucidated with reference to the embodiments described hereinafter.

In the drawings:

Fig. 1 is a side view, partly cut away and partly in cross-section, of an electric lamp in accordance with the invention comprising a lamp cap;

Fig. 2 shows an electric lamp provided with a reflector and an adapter;

Fig. 3 shows, in a part of a C.I.E. 1931 color triangle diagram, the color coordinates of an electric lamp comprising a coating in accordance with the invention.

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The Figures are purely schematic and not drawn to scale. Particularly for clarity, some dimensions are exaggerated strongly. In the Figures, like reference numerals refer to like parts whenever possible.

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Fig. 1 shows an electric lamp in accordance with the invention, a part of which is shown in side view, partly cut away, and another part of which is shown in cross-section. The electric lamp comprises a light-transmitting lamp vessel 1, for example made of glass, which is closed in a gastight manner and in which an electric element 2, being a (spiral-shaped) tungsten incandescent body with a center 4 in the Figure, is axially positioned on an axis 5 and is connected to current conductors 6 which issue from the lamp vessel to the exterior. The lamp shown has a filling of an inert gas, for example an Ar/Ne mixture, with a filling pressure slightly above 5 bar.

A lamp cap 10 is firmly connected to the lamp vessel 1. The lamp cap 10 has a synthetic resin housing 11. The housing 11 comprises a flat base portion 7 which is at least substantially perpendicular to the axis 5. The lamp vessel 1 is closed off in a gastight manner by means of a plate 8 of an insulating material, which plate lies in a plane which is at least substantially perpendicular to the axis 5. Electric element 2 is mounted in a previously defined position with respect to the plate 8 during the manufacture of the lamp. The plate 8 of the lamp vessel 1 is pressed home against the base portion by locking means 9, for example ridges, such that the electric element 2 will enter a previously defined position with respect to the reference means 12, for example studs. The studs 12 form part of the lamp cap and are designed to abut against a support 30, for example a reflector, as is visible in Fig. 2.

The lamp cap also comprises contact members 14 which are provided with a screen 13 and to which the current conductors 6 of the lamp vessel 1 are connected. A resilient intermediate portion 15, which is provided with coupling means 17, resilient tags in the Figure designed for coupling the reflector to the lamp cap, forms an integral whole with the housing 11. The resilient action of the intermediate portion is obtained in that the intermediate portion is made so as to be hollow, so that no more than a wall remains as the intermediate portion, whereupon a major portion of the wall is removed by means of two grooves 18 which run perpendicularly to the axis 5. The remaining portion of the wall forms

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a bridge 19 which is rotated, near the next groove, through an angle of, for example, 180° about the axis 5.

The lamp vessel 1 of the electric lamp has a relatively small axial dimension of approximately 22 mm and is suitable for consuming a relatively high power of, for example, 5 to 25 W. The electric lamp has a service life of approximately 6000 hours in this case.

In accordance with the invention, at least a part of the lamp vessel 1 is covered with a light-absorbing coating 3 having an average thickness of 2-3 μ m.

Fig. 2 shows the electric lamp provided with a support 30, being a reflector with a transparent plate 33 in the drawing, as well as with an adapter 25. In this configuration of a lamp with an adapter and a reflector, where the reflector is provided with a rubber ring 31 retained in a groove 32, the rubber ring seals off the opening 26 between the lamp cap and the reflector in a gastight manner. The adapter is provided with standardized contact points 27 which are passed through the bottom plate 28 of the adapter in a gastight manner and are connected to contact members 14 of the lamp cap 10.

It is visible in the drawing that the lamp cap 10 falls substantially entirely within a cone 36 which has its apex 35 in the center 4 of the electric element 2 and has an apex half angle α of 25°. The light originating from the electric element 2 can reach the reflecting surface 34 substantially without obstruction and is reflected there at least substantially axially in the direction of the transparent plate 33.

Example 1

A quantity of 10 g ZnFe₂O₄ (particle size 70 nm) is dispersed in a 50/50% water/ethanol mixture, using "disperbyk 190" as the dispersing agent. The overall weight of the mixture is 30 g. By means of wet ball milling using 2 mm zirconium oxide grains, an optically clear liquid is obtained.

A quantity of 3 g Fe₂O₃ (particle size 40 nm) is dispersed in a corresponding manner.

A hydrolysis mixture of 40 g methyltrimethoxy silane (MTMS), 0.6 g tetraethylorthosilicate (TEOS), 32 g water, 4 g ethanol and 0.15 g glacial acetic acid is stirred for 48 hours at room temperature and, subsequently, stored in a refrigerator.

A coating liquid is prepared by mixing 10 g of said ZnFe₂O₄ dispersion, 6 g of the Fe₂O₃ dispersion and 10 g of the MTMS/TEOS hydrolysis mixture with 4 g methoxy propanol, which coating liquid is subsequently spray coated onto the outer surface of the

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major part of a lamp vessel. The coating is cured for 10 minutes at a temperature of 250°C. In this manner, a light-absorbing coating in a thickness up to 3 µm is obtained on a glass lamp vessel without crack formation during drying and curing.

An electric lamp provided with a light-absorbing coating manufactured as described in this embodiment, is amber-colored, transparent and free of light scattering.

For a light-absorbing coating thus prepared, the color co-ordinate (x; y) in accordance with the definition of the C.I.E. 1931 color triangle diagram is (0.589; 0.405) at an overall transmission of 52% after the electric lamp has burned for 1 hour at 350 °C. The color point of the coating is stable during the service life of the electric lamp.

The coating obtained in accordance with the recipe has a thickness of 2.7 μ m. The weight fraction of the components in this coating is 52% ZnFe₂O₄ and Fe₂O₃, 18% "disperbyk 190", and 30% MTMS.

Fig. 3 shows, in a part of a C.I.E. 1931 color triangle diagram, the color coordinates (x; y) of an electric lamp with a light-absorbing coating comprising the above-described mixture of ZnFe₂O₄ and Fe₂O₃ (indicated by means of circles in Fig. 3). The circle with the lowest x-coordinate corresponds to a layer thickness of the ZnFe₂O₄/Fe₂O₃ coating of approximately 2 μm. The circle with the highest x-coordinate corresponds to a layer thickness of the ZnFe₂O₄/Fe₂O₃ coating of approximately 3 μm. For comparison, the color point of a coating comprising only Fe₂O₃ is given for a range of layer thicknesses (triangles: variation in thickness from 1.5 to 3 μm in the hatched area). Fig. 3 also shows two specified areas in the color triangle, wherein the color point of an electric lamp used as an amber-colored indicator for an automobile should be situated. The hatched area referenced S1 corresponds to the European ECE standard for an amber-colored indicator, and the area referenced S2 corresponds to the American SAE standard for an amber-colored indicator, which standards are both known to those skilled in the art. The light-absorbing coating obtained which is applied to an electric lamp can suitably be used as an amber-colored indicator and passes the Fakra test known to those skilled in the art.

Example 2

A quantity of 3 g BiVO₄ is dispersed in a 50/50% water/ethanol mixture, using "solsperse 41090" as the dispersing agent. The overall weight of the mixture is 23 g. By means of wet ball milling using 2 mm zirconium oxide grains, a stable dispersion is obtained.

A quantity of 3 g Fe₂O₃ is dispersed in a corresponding manner.

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A hydrolysis mixture of 40 g methyltrimethoxy silane (MTMS), 0.6 g tetraethyl orthosilicate (TEOS), 32 g water, 4 g ethanol and 0.15 g glacial acetic acid is stirred for 48 hours at room temperature and subsequently stored in a refrigerator.

A coating liquid is prepared by mixing 10 g of said BiVO₄ dispersion, 6 g of the Fe₂O₃ dispersion and 10 g of the MTMS/TEOS hydrolysis mixture with 4 g methoxy propanol, whereafter the coating liquid is spray coated onto the outer surface of the major part of a lamp vessel. The coating is dried for 20 minutes at a temperature of 160°C. In this manner, a light-absorbing coating in a thickness up to 3 μ m is formed on a glass lamp vessel without crack formation during drying and curing.

An electric lamp provided with a light-absorbing coating made in accordance with the embodiment described herein is amber-colored and relatively free of light scattering, although the diameter of the bismuth-vanadate particles exceeds 100 nm.

After the electric lamp has burned for one hour, the color co-ordinate (x; y) for a light-absorbing coating thus prepared amounts to (0.592; 0.101), in accordance with the definition of the C.I.E. 1931 color triangle diagram, at an overall transmission of 50%. The color point of the coating remains stable during the service life of the electric lamp.

The coating obtained in accordance with the recipe has a thickness of at least substantially 3 μm . The weight fraction of the components in this coating is 21% Fe₂O₃, 21% BiVO₄, 17% solspers and 41% MTMS.

Fig. 3 shows the color co-ordinates (x; y) of an electric lamp with a light-absorbing coating comprising the above-described mixture of BiVO₄ and Fe₂O₃ (indicated by means of the squares in Fig. 3). The square with the lowest x-coordinate corresponds to a layer thickness of the BiVO₄/Fe₂O₃ coating of approximately 2 μ m. The square with the highest x-coordinate corresponds to a layer thickness of the BiVO₄/Fe₂O₃ coating of approximately 3 μ m.

Example 3

A quantity of 6 g P-doped Fe₂O₃ is dispersed in a 50/50% water/ethanol mixture, using "disperbyk 190" as the dispersing agent. The overall weight of the mixture is 32 g.

A hydrolysis mixture of 40 g methyltrimethoxy silane (MTMS), 0.6 g tetraethyl orthosilicate (TEOS), 32 g water, 4 g ethanol and 0.15 g glacial acetic acid is stirred for 48 hours at room temperature and subsequently stored in a refrigerator.

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A coating is prepared by mixing 20 g of the P-doped Fe₂O₃ dispersion and 7 g of the MTMS/TEOS hydrolysis mixture with 8 g methoxy propanol, and said coating liquid is subsequently spray coated onto the outer surface of the major part of a lamp vessel. The coating is dried for 20 minutes at a temperature of 160 °C. In this manner, a light-absorbing coating having a thickness up to 6 μ m is formed on a glass lamp vessel without crack formation during drying and curing. The realization of such a relatively large layer thickness is possible because a relatively high concentration of pigment is applied at a relatively low concentration of MTMS.

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An electric lamp provided with a light-absorbing coating manufactured in accordance with the embodiment described herein is red, transparent and free of light scattering.

After the electric lamp has burned for one hour, the color co-ordinate (x; y) for a light-absorbing coating thus prepared is (0.665; 0.335), in accordance with the definition of the C.I.E. 1931 color triangle diagram, at an overall transmission of approximately 20%. The color point of the coating remains stable throughout the service life of the electric lamp.

The specified color co-ordinates, including the color point of an electric lamp used as a red brake light for an automobile, are, in accordance with the European ECE standard known to those skilled in the art: x = 0.665, y = 0.335; x = 0.657, y = 0.335; x = 0.7307; y = 0.2613, and in accordance with the American NA-standard known to those skilled in the art: x = 0.67, y = 0.33; x = 0.65, y = 0.33; x = 0.71, y = 0.27. The color point of the electric lamp provided with a light-absorbing coating made in accordance with the embodiment described herein lies within the specified areas for red brake light.

It will be clear that, within the scope of the invention, many variations are possible to those skilled in the art. In the sol-gel process, many alternative preparation methods are possible. For example, for the acid used to hydrolyze use can alternatively be made of maleic acid. Furthermore, it is also possible to use pigment combinations to cause the color point to shift towards red. Besides, the color temperature of the light to be emitted by the electric lamp can be increased while, for example, the color co-ordinates remain substantially positioned on the black body locus.

The scope of protection of the invention is not limited to the examples given herein. The invention is embodied in each novel characteristic and each combination of characteristics. Reference numerals in the claims do not limit the scope of protection thereof. The use of the term "comprising" does not exclude the presence of elements other than those

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mentioned in the claims. The use of the word "a" or "an" before an element does not exclude the presence of a plurality of such elements.

CLAIMS:

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1. An electric lamp comprising a light-transmitting lamp vessel (1) which accommodates a light source (2),

wherein at least a part of the lamp vessel (2) is provided with a light-absorbing coating (3),

wherein the light-absorbing coating (3) includes a pigment which absorbs a part of the visible light,

characterized in that

the light-absorbing coating (3) comprises a network which can be obtained by conversion of an organically modified silane by means of a sol-gel process,

said organically modified silane being selected from the group formed by compounds of the following structural formula: R^ISi(OR^{II})₃,

wherein R^I comprises an alkyl group or an aryl group, and wherein R^{II} comprises an alkyl group.

- 2. An electric lamp as claimed in claim 1, characterized in that the R¹ group comprises CH₃ or C₆H₅.
 - 3. An electric lamp as claimed in claim 1 or 2, characterized in that the R^{II} group comprises CH_3 or C_2H_5 .
 - 4. An electric lamp as claimed in claim 1 or 2, characterized in that an average diameter d_p of the pigment is $d_p \le 100$ nm.
- 5. An electric lamp as claimed in claim 1 or 2, characterized in that the thickness
 25 t_c of the light-absorbing coating (3) is t_c ≥ 1 μm.
 - 6. An electric lamp as claimed in claim 1 or 2, characterized in that silica particles having a diameter $d \le 50$ nm are incorporated in the network.

7. An electric lamp as claimed in claim 1 or 2, characterized in that the pigment is an inorganic pigment.

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- 8. An electric lamp as claimed in claim 7, characterized in that the pigment is selected from the group formed by iron oxide, iron oxide doped with phosphor, zinc-iron oxide, cobalt aluminate, neodymium oxide, bismuth vanadate, zirconium praseodymium silicate or mixtures thereof.
- 9. An electric lamp as claimed in claim 8, characterized in that the pigment is
 10 formed by a mixture of iron oxide and bismuth vanadate or by a mixture of iron oxide doped with phosphor and bismuth vanadate.

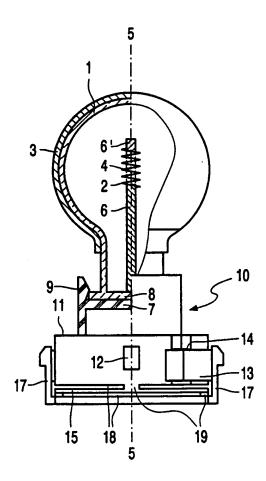
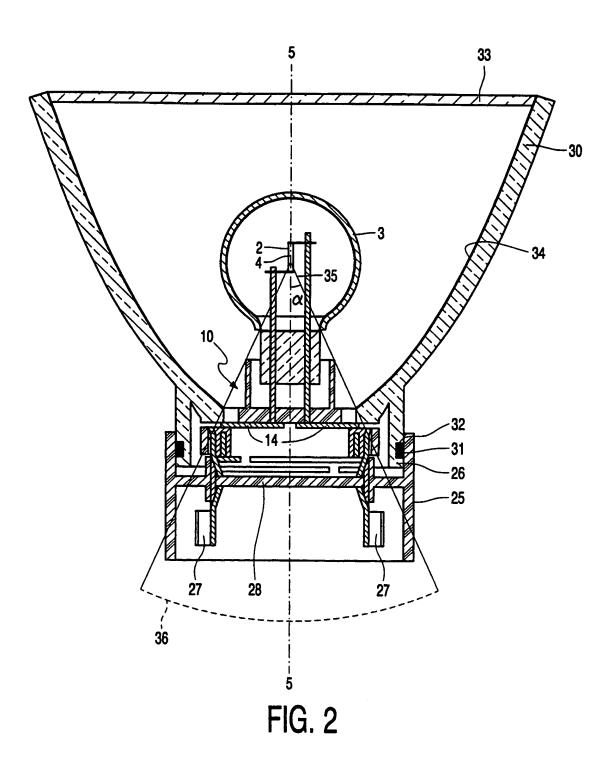
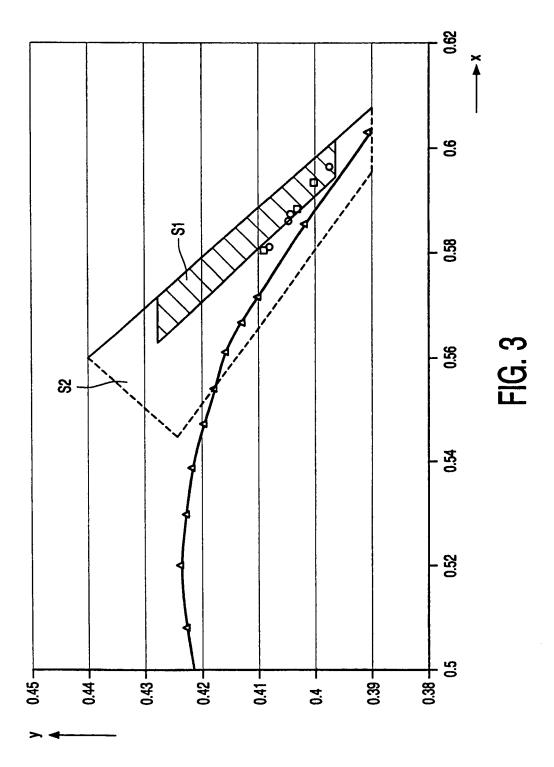


FIG. 1





INTERNATIONAL SEARCH REPORT

Interr nal Application No PCT/EP 00/08502

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 H01J61/40							
According to International Patent Classification (IPC) or to both national classification and IPC							
B. FIELDS	and the second s						
Minimum documentation searched (classification system followed by classification symbols) IPC 7 H01J H01K G02B							
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched							
Electronic da	ata base consulted during the international search (name of data base	e and, where practical, search terms used)				
	ternal, WPI Data, PAJ						
C. DOCUMENTS CONSIDERED TO BE RELEVANT							
Category °	Citation of document, with indication, where appropriate, of the rele	vant passages	Relevant to claim No.				
Α	DATABASE WPI Section Ch,, 1968		1-3				
	Derwent Publications Ltd., London Class A00, AN 1968-91495P XP002155448 & CA 766 196 A (WESTINGHOUSE ELEC CORPORATION), 3 March 1965 (1965- cited in the application abstract						
A	EP 0 874 278 A (AGFA GEVAERT NV) 28 October 1998 (1998-10-28) abstract page 6, line 39 - line 45 page 6, line 50 -page 7, line 4		1				
Furt	ther documents are listed in the continuation of box C.	Patent family members are listed	in annex.				
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Information on patent family members

Interr. nal Application No PCT/EP 00/08502

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
CA 766196	Α		NONE	
EP 0874278	Α	28-10-1998	NONE	